Source URL: https://medicines.health.europa.eu/veterinary/en/600000041736

M+PAC

Not authorised

• Mycoplasma hyopneumoniae, Inactivated

Product identification

Medicine name:

M+PAC >=1.47 RPU ENEΣΙΜΟ ΓΑΛΑΚΤΩΜΑ M+PAC

Active substance:

Mycoplasma hyopneumoniae, Inactivated

Target species:

Pig

Route of administration:

Intramuscular use

Product details

Active substance and strength:

Mycoplasma hyopneumoniae, Inactivated 1.47 relative unit(s) / 1.00 millilitre(s)

Pharmaceutical form:

Emulsion for injection

Withdrawal period by route of administration:

Intramuscular use:

Pig

- Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI09AB13

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Surrendered

Authorised in:

Greece

Package description:

Box of 10 bottle of 200 ml

Box of 5 bottle of 200 ml

Box of 2 bottle of 200 ml

Box of 1 bottle of 200 ml

Box of 1 bottle of 50 ml

Box of 10 bottle of 100 ml

Box of 5 bottle of 100 ml

Box of 2 bottle of 100 ml

Box of 1 bottle of 100 ml

Box of 10 bottles of 50 ml

Box of 5 bottles of 50 ml

Box of 2 bottles of 50 ml

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Immunological veterinary medicinal product application (Article 13d of Directive No 2001/82/EC)

Marketing authorisation holder:
Intervet Hellas A.E.

Marketing authorisation date:
10/10/2002

Manufacturing sites for batch release:
Burgwedel Biotech GmbH

Responsible authority:
National Organization For Medicines

Authorisation number:
24501/03-04-2009/K-0143801

Date of authorisation status change:
2/04/2009

Reference member state:

Hungary

Procedure number:

HU/V/0140/001/MR

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet